



DEPARTMENT OF JUSTICE
DRUG ENFORCEMENT ADMINISTRATION
MANUFACTURER OF CONTROLLED SUBSTANCES
NOTICE OF APPLICATION
GE HEALTHCARE

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on August 31, 2012, GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004-1412, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to manufacture a radioactive product to diagnose Parkinson's disease; and to manufacture a bulk investigational new drug (IND) for clinical trials.

Any other such applicant, and any person who is presently registered with DEA to manufacture such a substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement

Administration, Office of Diversion Control, Federal
Register Representative (ODL), 8701 Morrisette Drive,
Springfield, Virginia 22152; and must be filed no later
than [INSERT DATE 60 DAYS FROM DATE OF PUBLICATION].

Joseph T. Rannazzisi
Deputy Assistant Administrator
Office of Diversion Control
Drug Enforcement Administration

DATED: February 8, 2013

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